This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Presently amended) A compound having the formula

$$R^{1}$$
 R^{2}
 $N-R_{5}$
 R^{1}
 $N-R^{3}$
 R^{6}
 R^{1}
 R^{6}
 R^{7}
 R^{7}
 R^{7}
 R^{7}
 R^{7}
 R^{7}

wherein:

 X^1 and X^2 are independently a direct bond or a linking atom or group selected from the group consisting of $-C(=X^3)-$, $-C(=X^3)-$ N(R^8)- $C(=X^3)-$;

 X^3 is -O- or -S-;

 R^1 is acyl of from about 7 $\underline{16}$ to about 23 carbons;

R² is hydrogen or lower alkyl;

R³ is alkylene of from 1 to about 10 carbons;

R⁴ is acyl of from about 7 16 to about 23 carbons;

R⁵ is hydrogen or lower alkyl;

R⁶ is a direct bond;

R⁷ is a direct bond or alkylene of from 1 to about 10 carbons;

R⁸ is hydrogen or lower alkyl;

P is a hydrophilic polymer; and

T is a targeting ligand which targets cells or receptors selected from the group consisting of myocardial cells, endothelial cells, epithelial cells, tumor cells and the glycoprotein GPIIbIIIa receptor.

2. Cancelled.

Office Action Dated: January 14, 2004

(Original) A compound according to Claim 2 wherein: 3.

 X^{1} is -C(=O)-NH-C(=O)-;

 X^2 is -C(=O)-;

R¹ is acyl of from about 15 to about 20 carbons;

R³ is alkylene of from 1 to about 3 carbons;

R4 is acyl of from about 15 to about 20 carbons; and

R⁶ is a direct bond;

R⁷ is lower alkylene.

(Original) A compound according to Claim 3 wherein: 4.

R¹ is acyl of from about 17 to about 19 carbons;

R³ is methylene;

R⁴ is acyl of from about 17 to about 19 carbons; and

R⁷ is ethylene.

- Cancelled. 5.
- (Original) A compound according to Claim 1 wherein said hydrophilic 6. polymer is selected from the group consisting of polyalkyleneoxides, polyvinyl alcohol, polyvinylpyrrolidones, polyacrylamides, polymethacrylamides, polyphosphazenes, poly(hydroxyalkylcarboxylic acids) and polyoxazolidines.
- (Original) A compound according to Claim 6 wherein said hydrophilic 7. polymer comprises a polyalkyleneoxide.
- (Original) A compound according to Claim 7 wherein said hydrophilic 8. polymer is selected from the group consisting of polyethylene glycol and polypropylene glycol.
- (Original) A compound according to Claim 8 wherein said hydrophilic 9. polymer is polyethylene glycol.

- (Original) A compound according to Claim 8 wherein said hydrophilic 10. polymer is PEG3400.
- (Original) A compound according to Claim 1 wherein said targeting 11. ligand comprises a peptide of the formula:

wherein:

m and n are independently an integer of from 1 to about 100;

Xaa and Zaa are independently selected from the group consisting of natural amino acids and synthetic amino acids;

Yaa is selected from Arginine, Homoarginine, and Lysine-N-

acetimidate; and

with the proviso that when Xaa and Zaa are sulfur containing amino acids, Xaa and Zaa may be linked together via a disulfide linkage.

> (Withdrawn) A compound according to Claim 11, wherein: 12.

> > Xaa is Glycine;

Yaa is Arginine;

Zaa is Serine;

n is 1, 2 or 3; and

m is 1.

- (Withdrawn) A compound according to Claim 12, wherein: 13. n is 3.
- (Original) A compound according to Claim 11, wherein: 14. Xaa and Zaa comprise an amino acid independently selected from sulfur containing amino acids.

DOCKET NO.: UNGR-1598 **Application No.:** 09/699,679

Office Action Dated: January 14, 2004

PATENT REPLY FILED UNDER EXPEDITED PROCEDURE PURSUANT TO 37 CFR § 1.116

15. (Original) A compound according to Claim 1 wherein said targeting ligand comprises a peptide of the following formula:

$$S \longrightarrow S$$
 $S \longrightarrow S$ $(Xaa)_x$ -Saa- $(Xaa)_x$ -Yaa-Gly-Asp- $(Zaa)_y$ -Saa- $(Zaa)_y$

wherein:

each x and y is independently an integer of from 0 to about 50; each Saa is selected from the group consisting of natural and synthetic sulfur containing amino acids;

each Xaa and Zaa are independently selected from the group consisting of natural amino acids and synthetic amino acids; and

Yaa is selected from Arginine, Homoarginine, and Lysine-Nacetimidate.

- 16. (Original) A compound according to Claim 15 wherein:
 each Saa is independently selected from the group consisting of DCysteine, L- Cysteine, D-Penicillamine and L-Penicillamine.
- 17. (Original) A targeted vesicle composition for therapeutic or diagnostic use *in vivo* comprising, in an aqueous carrier, lipid, protein or polymer gas filled vesicles, wherein said vesicles further comprise a compound according to Claim 1.
- 18. (Original) A targeted vesicle composition according to Claim 17, wherein said vesicles are selected from the group consisting of liposomes and micelles.
- 19. (Original) A targeted vesicle composition according to Claim 18, wherein said vesicles comprise liposomes.

DOCKET NO.: UNGR-1598 **Application No.:** 09/699,679

Office Action Dated: January 14, 2004

PATENT REPLY FILED UNDER EXPEDITED PROCEDURE PURSUANT TO 37 CFR § 1.116

- 20. (Original) A targeted vesicle composition according to Claim 19 wherein said liposomes comprise a phospholipid selected from the group consisting of phosphatidylcholine, phosphatidylethanolamine and phosphatidic acid.
- 21. (Original) A targeted vesicle composition according to Claim 20 wherein said phosphatidylcholine is selected from the group consisting of dioleoylphosphatidyl-choline, dimyristoylphosphatidylcholine, dipalmitoylphosphatidylcholine and distearoylphosphatidylcholine.
- 22. (Original) A targeted vesicle composition according to Claim 21 wherein said phosphatidylcholine comprises dipalmitoylphosphatidylcholine.
- 23. (Original) A targeted vesicle composition according to Claim 20 wherein said phosphatidylethanolamine is selected from the group consisting of dipalmitoyl-phosphatidylethanolamine, dioleoylphosphatidylethanolamine, N-succinyldioleoyl-phosphatidylethanolamine and 1-hexadecyl-2-palmitoylglycerophosphoethanolamine.
- 24. (Original) A targeted vesicle composition according to Claim 23 wherein said phosphatidylethanolamine comprises dipalmitoylphosphatidylethanolamine.
- 25. (Original) A targeted vesicle composition according to Claim 20 wherein said phosphatidic acid comprises dipalmitoylphosphatidic acid.
- 26. (Original) A targeted vesicle composition according to Claim 17, wherein said vesicles comprise a gas selected from the group consisting of perfluorocarbons and sulfur hexafluoride.
- 27. (Original) A targeted vesicle composition according to Claim 26 wherein said perfluorocarbon gas is selected from the group consisting of perfluoromethane, perfluoropropane, perfluorobutane and perfluorocyclobutane.

PATENT REPLY FILED UNDER EXPEDITED PROCEDURE PURSUANT TO 37 CFR § 1.116

- 28. (Original) A targeted vesicle composition according to Claim 27 wherein said perfluorocarbon gas is selected from the group consisting of perfluoropropane and perfluorobutane.
- 29. (Original) A targeted vesicle composition according to Claim 28 wherein said perfluorocarbon gas comprises perfluorobutane.
- 30. (Original) A targeted vesicle composition according to Claim 17 wherein said gas is derived, at least in part, from a gaseous precursor.
- 31. (Original) A targeted vesicle composition according to Claim 30 wherein said gaseous precursor has a boiling point of greater than about 37°C.
- 32. (Original) A targeted vesicle composition according to Claim 31 wherein said gaseous precursor comprises a perfluorocarbon.
- 33. (Original) A targeted vesicle composition according to Claim 32 wherein said perfluorocarbon is selected from the group consisting of perfluoropentane and perfluorohexane.
- 34. (Original) A targeted vesicle composition according to Claim 17 wherein said vesicles further comprise a bioactive agent that is different from said gas and said compound.
- 35. (Original) A targeted vesicle composition according to Claim 34 wherein said bioactive agent comprises a therapeutic agent selected from the group consisting of genetic material, dihydroergotamine, heparin sulfate, tissue plasminogen activator, streptokinase, urokinase, hirudin, and mixtures thereof.
 - 36-53. Cancelled.

Office Action Dated: January 14, 2004

37 CFR § 1.116

54. (Previously presented) A compound according to Claim 1, wherein:

$$X^1$$
 is $-C(=X^3)-N(R^8)-$;

$$X^2$$
 is $C(=X^3)$;

$$X^3$$
 is O;

R¹ is acyl of 18 carbons;

 R^2 is H:

R³ is ethylene;

R⁴ is acyl of 18 carbons;

R⁵ is H;

R⁶ is a direct bond;

R⁷ is ethylene;

 R^8 is H;

P is PEG-3400; and

T comprises a peptide having the sequence CRGDC, wherein the two cysteines are linked together via a disulfide linkage.

- 55. (Previously presented) A targeted vesicle composition for therapeutic or diagnostic use *in vivo* comprising, in an aqueous carrier, lipid vesicles, wherein said vesicles comprise a compound according to Claim 54.
- 56. (Previously presented) A targeted vesicle composition according to Claim 55 wherein said lipid vesicles comprise a phospholipid selected from the group consisting of phosphatidylcholine, phosphatidylethanolamine and phosphatidic acid.

- 57. (Previously presented) A targeted vesicle composition according to Claim 56 wherein said phosphatidylethanolamine comprises dipalmitoylphosphatidylethanolamine.
- 58. (Previously presented) A targeted vesicle composition according to Claim 55, wherein said vesicles comprise a gas selected from the group consisting of perfluorocarbons and sulfur hexafluoride.
- 59. (Previously presented) A targeted vesicle composition according to Claim 58, wherein said vesicles comprise perfluorobutane.
- 60. (Previously presented) A targeted vesicle composition according to Claim 55, further comprising urokinase.
 - 61. (New) A compound according to Claim 4 wherein: $R^1 \text{ and } R^4 \text{ are acyl of about } 18 \text{ carbons.}$
 - (New) A compound according to Claim 4 wherein:R¹ is an acyl of from about 17 to about 19 carbons.
 - (New) A compound according to Claim 4 wherein:R¹ is an acyl of about 18 carbons.